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DATE: Saturday, September 17, 2005

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DB=DWPI; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L28	L27 and l26	62
<input type="checkbox"/>	L27	toxin	7426
<input type="checkbox"/>	L26	quinquestriatus or leiurus or scorpion	652

DB=JPAB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L25	quinquestriatus or leiurus or scorpion	6
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DB=EPAB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L24	L23 and l22	9
<input type="checkbox"/>	L23	toxin	1177
<input type="checkbox"/>	L22	quinquestriatus or leiurus or scorpion	29

DB=USOC; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L21	L20 same l19	0
<input type="checkbox"/>	L20	toxin	463
<input type="checkbox"/>	L19	quinquestriatus or leiurus or scorpion	28
<input type="checkbox"/>	L18	potassium or k+	105038

DB=PGPB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L17	L15 and (potassium or k+)	109
<input type="checkbox"/>	L15	(L14 or l12) same l13	171
<input type="checkbox"/>	L14	scorpion	835
<input type="checkbox"/>	L13	toxin	22638
<input type="checkbox"/>	L12	quinquestriatus or leiurus	43

DB=USPT; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L11	l7 and l6 not l9	61
<input type="checkbox"/>	L10	l7 and l3 not l8	33
<input type="checkbox"/>	L9	l7 same l6	34
<input type="checkbox"/>	L8	l7 same l3	3
<input type="checkbox"/>	L7	potassium or k+	282322
<input type="checkbox"/>	L6	L5 not l3	208
<input type="checkbox"/>	L5	l4 same l2	258
<input type="checkbox"/>	L4	scorpion	1172
<input type="checkbox"/>	L3	l1 same L2	53
<input type="checkbox"/>	L2	toxin	22671

L1 quinquestriatus or leiurus

79

END OF SEARCH HISTORY

S39	S38 AND S2	E30	1 AU=LEE JK-Y	Ref	Items Index-term
S40	338 ID (sorted in duplicate order)	E31	77 AU=LEE JL	E1	15 AU=LEE JIAHN-SHING
S41	3201080 DNA OR GENE OR PLASMID	E32	4 AU=LEE JLC	E2	2 AU=LEE JIAHN-TE
S42	33 S39 AND S41	E33	2 AU=LEE JLJR	E3	0 AU=LEE JIAN
S43	33 ID (sorted in duplicate order)	E34	1 AU=LEE JL-C	E4	1 AU=LEE JIAN TAO
		E35	1005 AU=LEE JM	E5	1 AU=LEE JIAN-CHENG
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		E37	7 AU=LEE JMH	E7	13 AU=LEE JIAN-MING
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		E39	1 AU=LEE JM JEANETTE	E9	1 AU=LEE JIAN GWU
		E40	3 AU=LEE JM JR	E10	4 AU=LEE JIAN-N-DER
		E41	1 AU=LEE JM L	E11	5 AU=LEE JIANN-FENG
		E42	1 AU=LEE JM TAMBLYN	E12	1 AU=LEE JIANN-GWU
		E43	2 AU=LEE JM-F	E13	1 AU=LEE JIANN-HSIUNG
		E44	2 AU=LEE J MARSHALL	E14	3 AU=LEE JIANN-SHU
		E45	1 AU=LEE J MARTIN		
		E46	1 AU=LEE J MI		
		E47	26 AU=LEE J MICHAEL		
		E48	162 AU=LEE J N		
				S46	13 AU="LEE JIAN-MING"
				S47	1034 S46 OR S45
				S48	3 S47 AND (S1 or S17)
S44	4131 AU="LEE J"	S45	1021 E35-E38,E41,E43		
		6/6/1	(Item 1 from file: 5) 0007115178 BIOSIS NO.: 199089033069 ANALYSIS OF THE BLOCKING ACTIVITY OF CHARYBDOTOXIN HOMOLOGS AND IODINATED DERIVATIVES AGAINST CALCIUM-ACTIVATED POTASSIUM CHANNELS 1989	6/6/15	(Item 15 from file: 5) 0006734218 BIOSIS NO.: 198784088367 CHARYBDOTOXIN SELECTIVELY BLOCKS SMALL CALCIUM-ACTIVATED POTASSIUM CHANNELS IN APLYSIA NEURONS 1987
		6/6/2	(Item 2 from file: 15) 08751880 PMID: 2477548 Analysis of the blocking activity of charybdotoxin homologs and iodinated derivatives against Ca^{2+} -activated K^+ -channels. Aug 1989	6/6/16	(Item 16 from file: 5) 0007723137 BIOSIS NO.: 19911968028 CHARYBDOTOXIN-SENSITIVE CALCIUM ACTIVATED POTASSIUM CHANNEL IS NOT INVOLVED IN GLUCOSE-INDUCED ELECTRICAL ACTIVITY IN PANCREATIC BETA-CELLS 1991
		6/6/3	(Item 3 from file: 5) 0014612756 BIOSIS NO.: 2003010581475 Atypical relaxation by scorpion venom in the lamb urethral smooth muscle involves both NO-dependent and -independent responses. 2003	6/6/17	(Item 17 from file: 155) 09417323 PMID: 1710672 Charybdotoxin -sensitive $K(Ca)$ channel is not involved in glucose-induced electrical activity in pancreatic beta-cells. Jan 1991
		6/6/4	(Item 4 from file: 5) 0010224284 BIOSIS NO.: 199698692117 Characterization of Ca^{2+} -activated 86RB $^{+}$ fluxes in rat C6 glioma cells: A system for identifying novel $IK\alpha$ -channel toxins 1986	6/6/18	(Item 18 from file: 5) 0011343339 BIOSIS NO.: 199800143536 Consequence of the removal of evolutionary conserved disulfide bridges on particular cysteine spacing govern specific disulfide bond formation 1988
		6/6/5	(Item 5 from file: 155) 115090117 PMID: 8821537 Characterization of Ca^{2+} -activated 86RB $^{+}$ fluxes in rat C6 glioma cells: a system for identifying novel $IK\alpha$ -channel toxins. Feb 1996	6/6/19	(Item 19 from file: 5) 0007735556 BIOSIS NO.: 19911918447 DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXIN A PEPTIDE BLOCKER OF POTASSIUM ION CHANNELS 1991
		6/6/6	(Item 6 from file: 155) 08675595 PMID: 2473920 Charybdotoxin blocks both Ca -activated K channels and Ca -independent voltage-gated K channels in rat brain synaptosomes. Jul 3 1989	6/6/20	(Item 20 from file: 155) 08571405 PMID: 2468961 Effect of some potassium channel blockers on contractile responses of the rabbit aorta. Feb 1989
		6/6/7	(Item 7 from file: 155) 08710107 PMID: 2475579 Charybdotoxin blocks voltage-gated K $^{+}$ channels in human and murine T lymphocytes. Jun 1989	6/6/21	(Item 21 from file: 5) 0006633771 BIOSIS NO.: 198987056652 EFFECT OF SOME POTASSIUM CHANNEL BLOCKERS ON CONTRACTILE RESPONSES OF THE RABBIT AORTA 1989
		6/6/8	(Item 8 from file: 5) 0006742983 BIOSIS NO.: 198988056098 CHARYBDOTOXIN BLOCKS VOLTAGE-GATED POTASSIUM CHANNELS IN HUMAN AND MURINE T LYMPHOCYTES 1989	6/6/22	(Item 22 from file: 155) 08812992 PMID: 2531622 Effects of potassium channel toxins from <i>Leiurus quinquestrigatus hebraeus</i> venom on responses to cromakalim in rabbit blood vessels. Nov 1989
		6/6/9	(Item 9 from file: 155) 11058434 PMID: 7543240 Charybdotoxin and its effects on potassium channels. Jul 1995	6/6/24	(Item 24 from file: 155) 10239832 PMID: 7667466 Influence of protein surface charge on the bimolecular kinetics of a potassium channel peptide inhibitor. Jul 13 1993
		6/6/10	(Item 10 from file: 5) 0009978314 BIOSIS NO.: 199598446147 Charybdotoxin and its effects on potassium channels [95]	6/6/23	(Item 23 from file: 5) 0007102492 BIOSIS NO.: 199089020333 EFFECTS OF POTASSIUM CHANNEL TOXINS FROM <i>LEIURUS-QUINQUESTRIGATUS-HEBRAEUS</i> VENOM ON RESPONSES TO CROMAKALIM IN RABBIT BLOOD VESSELS 1989
		6/6/11	(Item 11 from file: 155) 08659199 PMID: 2482078 Charybdotoxin is a new member of the K $^{+}$ -channel toxin family that includes dendrotoxin I and mast cell degranulating peptide. Dec 12 1989	6/6/25	(Item 25 from file: 5) 0009404393 BIOSIS NO.: 199497425678 Evidence in support of a role for Ca^{2+} -activated K $^{+}$ channels in the hamster sperm acrosome reaction 1994
		6/6/12	(Item 12 from file: 5) 000713033 BIOSIS NO.: 199089050924 CHARYBDOTOXIN IS A NEW MEMBER OF THE POTASSIUM CHANNEL TOXIN FAMILY THAT INCLUDES DENDROTOXIN I AND MAST CELL DEGRANULATING PEPTIDE 1989	6/6/26	(Item 26 from file: 155) 10570679 PMID: 7520055 Evidence in support of a role for Ca^{2+} -activated K $^{+}$ channels in the hamster sperm acrosome reaction 1994
		6/6/13	(Item 13 from file: 5) 0006941512 BIOSIS NO.: 199038119403 CHARYBDOTOXIN IS A NEW MEMBER OF THE TOXIN FAMILY THAT INCLUDES DENDROTOXIN I AND MCD AND BLOCKS DENDROTOXIN-SENSITIVE VOLTAGE-ACTIVATED POTASSIUM CHANNELS 1990	6/6/27	(Item 27 from file: 5) 0014215910 BIOSIS NO.: 200300174629 Functional analysis of an archaeabacterial voltage-dependent K $^{+}$ channel. 2003
		6/6/14	(Item 14 from file: 155) 07976191 PMID: 2442295 Charybdotoxin selectively blocks small Ca -activated K channels in Aplysia neurons. Jul 1987		

6/6/28 (Item 28 from file: 5) 0006267342 BIOSIS NO.: 1988861072633 PHARMACOLOGY OF POTASSIUM CHANNELS IN THE PLASMALEMMA OF THE GREEN ALGA CHARA-CORALLINA 1988

6/6/29 (Item 29 from file: 155) 07773735 PMID: 2433153 Identification of two toxins from scorpion (*Leurus quinquestratus*) venom which block distinct classes of calcium-activated potassium channel Dec 1 1986

6/6/30 (Item 30 from file: 155) 08722670 PMID: 2476127 Interactions between dendrotoxin, a blocker of voltage-dependent potassium channels, and charybdotoxin, a blocker of calcium-activated potassium channels, at binding sites on neuronal membranes. Aug 30 1989

6/6/31 (Item 31 from file: 5) 0006799478 BIOSIS NO.: 198988114593 INTERACTION BETWEEN DENDROTOXIN A BLOCKER OF VOLTAGE-DEPENDENT POTASSIUM CHANNELS AND CHARYBDOTOXIN A BLOCKER OF CALCIUM-ACTIVATED POTASSIUM CHANNELS AT BINDING SITES ON NEURONAL MEMBRANES 1989

6/6/32 (Item 32 from file: 5) 001988561 BIOSIS NO.: 200400359350 Kbot1, a three disulfide bridges toxin from *Buthus occitanus tunetanus* venom highly active on both SK and Kv channels 2004

6/6/33 (Item 33 from file: 5) 0006395808 BIOSIS NO.: 198926104699 LEURUS-QUINQUESTRATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES 1989

6/6/34 (Item 34 from file: 155) 10823421 PMID: 7819188 NMR sequential assignments and solution structure of charybdotoxin, a small scorpion toxin that blocks chloride channels. Jan 10 1995

6/6/35 (Item 35 from file: 5) 0006488287 BIOSIS NO.: 198598116660 NMR Sequential Assignments and Solution Structure of Charybdotoxin, a Small Scorpion Toxin That Blocks Chloride Channels 1995

6/6/36 (Item 36 from file: 155) 10805806 PMID: 7533951 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestratus hebreus*. Nov 1994

6/6/37 (Item 37 from file: 5) 0008575215 BIOSIS NO.: 198598043048 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestratus hebreus* 1994

6/6/38 (Item 38 from file: 155) 09990086 PMID: 1280139 Neurolustoxin and leurotoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels Sep 1992

6/6/39 (Item 39 from file: 5) 0008709107 BIOSIS NO.: 198395011373 Neurolustoxin and leurotoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels Sep 1992

6/6/40 (Item 40 from file: 155) 06794986 PMID: 6197125 A study on the membrane depolarization of skeletal muscles caused by a scorpion toxin, sea anemone toxin II and crotamine and the interaction between toxins. Jul 1983

6/6/41 (Item 41 from file: 5) 0007309715 BIOSIS NO.: 198900094194 POLARIZED RUBIDIUM-86 EFFLUXES IN PRIMARY CULTURES OF RABBIT KIDNEY PROXIMAL CELLS ROLE OF CALCIUM AND HYPOTONICITY 1990

6/6/42 (Item 42 from file: 155) 09066202 PMID: 2165808 Polarized 86Rb+ effluxes in primary cultures of rabbit kidney proximal cells: role of calcium and hypotonicity. Jul 9 1990

6/6/43 (Item 43 from file: 5) 0005560519 BIOSIS NO.: 198783029410 PURIFICATION OF CHARYBDOTOXIN A SPECIFIC INHIBITOR OF THE HIGH-CONDUCTANCE CALCIUM-ACTIVATED POTASSIUM CHANNEL 1986

6/6/44 (Item 44 from file: 155) 07702384 PMID: 2429598 Purification of charybdotoxin, a specific inhibitor of the high-conductance Ca2+-activated K+ channel Nov 5 1986

6/6/45 (Item 45 from file: 155) 08257702 PMID: 2453055 Purification, sequencing, and model structure of charybdotoxin, a potent selective inhibitor of calcium-activated potassium channels. May 1988

6/6/46 (Item 46 from file: 5) 0006204047 BIOSIS NO.: 198886043968 PURIFICATION SEQUENCE AND MODEL STRUCTURE OF CHARYBDOTOXIN A POTENT SELECTIVE INHIBITOR OF CALCIUM-ACTIVATED POTASSIUM CHANNELS 1986

6/6/47 (Item 47 from file: 5) 0009198685 BIOSIS NO.: 19949718150 Solution structure of a core peptide derived from scyllatoxin 1994

6/6/48 (Item 48 from file: 155) 13306597 PMID: 10831954 Solution structure of potassium channel-inhibiting scorpion toxin Lq2. Mar 1 1999

6/6/49 (Item 49 from file: 5) 0008984542 BIOSIS NO.: 199497005827 Synthesis of charybdotoxin and of two N-terminal truncated analogues: Structural and functional characterization 1993

6/6/50 (Item 50 from file: 5) 0007639197 BIOSIS NO.: 19919102688 SYNTHESIS AND STRUCTURAL CHARACTERIZATION OF CHARYBDOTOXIN A POTENT PEPTIDYL INHIBITOR OF THE HIGH CONDUCTANCE CALCIUM-ACTIVATED POTASSIUM ION CHANNEL 1990

6/6/51 (Item 51 from file: 155) 09151531 PMID: 1639936 Synthesis and structural characterization of charybdotoxin, a potent peptidyl inhibitor of the high conductance Ca2+-activated K+ channel Nov 5 1990

6/6/52 (Item 52 from file: 5) 0010206293 BIOSIS NO.: 199698674126 Synthesis and structural characterization of arabyotoxin, a potassium channel blocker charybdotoxin 1996

6/6/53 (Item 53 from file: 5) 0009025688 BIOSIS NO.: 199497046373 Toxin pharmacology of the large-conductance Ca2+-activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture 1993

6/6/54 (Item 54 from file: 155) 10398153 PMID: 75056914 Toxin pharmacology of the large-conductance Ca2+-activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture. Oct 1993

6/6/55 (Item 55 from file: 155) 0974404035 PMID: 1373656 Toxin pharmacology of the ATP-induced hyperpolarization in Madin-Darby canine kidney cells. Mar 23 1992

6/6/56 (Item 56 from file: 5) 0008332816 BIOSIS NO.: 199294034657 TOXIN PHARMACOLOGY OF ATP-INDUCED HYPERPOLARIZATION IN MADIN-DARBY CANINE KIDNEY CELLS 1992

6/6/57 (Item 57 from file: 5) 0006361782 BIOSIS NO.: 1989360906173 TOXINS IN THE CHARACTERIZATION OF POTASSIUM CHANNELS 1989

6/6/58 (Item 58 from file: 155) 10066261 PMID: 7678959 Toxin sensitivity of the calcium-dependent rubidium efflux in Madin-Darby canine kidney cells. Jan 29 1993

6/6/59 (Item 59 from file: 5) 0008801098 BIOSIS NO.: 199395103364 Toxin sensitivity of the calcium-dependent rubidium efflux in Madin-Darby canine kidney cells 1993

6/6/60 (Item 60 from file: 155) 08833309 PMID: 2600838 A voltage-dependent outward current with fast kinetics in single smooth muscle cells isolated from rabbit portal vein. May 1989

6/6/61 (Item 61 from file: 5) 0006708256 BIOSIS NO.: 198988023371 A VOLTAGE-DEPENDENT OUTWARD CURRENT WITH FAST KINETICS IN SINGLE SMOOTH MUSCLE CELLS ISOLATED FROM RABBIT PORTAL VEIN 1989

6/6/62 (Item 62 from file: 5) 0007742476 BIOSIS NO.: 19919125367 THREE-DIMENSIONAL STRUCTURE OF NATURAL CHARYBDOTOXIN IN AQUEOUS SOLUTION BY PROTON NMR CHARYBDOTOXIN POSSESSES A STRUCTURAL MOTIF FOUND IN OTHER SCORPION TOXINS 1991

6/7/2 (Item 2 from file: 155) DIALOG(R)File 155: MEDLINE(R) (C) format only 2005 Dialog. All its. reserv.

08751880 PMID: 2477548 Analysis of the blocking activity of charybdotoxin homologs and iodinated derivatives against Ca2+-activated K+ channels.

Lucchesi K, Ravindran A, Young H, Moczydlowski E Department of Pharmacology, Yale University School of Medicine, New Haven, Connecticut 06510. Journal of membrane biology (UNITED STATES) Aug 1989, 109 (3) p269-81, ISSN 0022-2631 Journal Code: 0211301 Contract/Grant No.: AR38796; AR; NIAMS; HL38156; HL; NHBL Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE, Completed

Two charybdotoxin peptides were purified from venom of the Israeli scorpion, *Leiurus quinquestratus hebraeus*. Microsequencing of the most abundant toxin, ChTX-Lq1, revealed identity with the 31-residue peptide previously sequenced by Gimenez-Gallego et al. [Gimenez-Gallego, G., et al., Proc. Natl. Acad. Sci. USA 85:3329-3333 (1988)]. Sequence data on the minor peptide, ChTX-Lq2, showed substantial homology to ChTX-Lq1 with differences observed at eight positions. These two charybdotoxin sequences, along with that of noxiustoxin, define a distinct family of scorpion peptide toxins with activity against K+ channels. Both charybdotoxin homologs inhibited Ca2+-dependent K+ efflux from human erythrocytes with similar potency, K0.5 approximately 40 nM. In planar bilayer assays of single K(Ca) channels from rat muscle, ChTX-Lq1 and ChTX-Lq2 blocked with intrinsic Kds of 1.3 and 43 nM, respectively, in the presence of 50 mM external KCl. A new application of dwell-time histogram analysis of single-channel blocking events was used to characterize the kinetic homogeneity of toxin samples and the blocking affinity of ChTX-Lq2 3

was the combined result of a faster dissociation rate and a slower association rate as compared to ChTX-Lq1. The blocking activity of two mono-iodinated derivatives of ChTX-Lq1 was also analyzed. Blocked dwell-time histograms of the iodinated peptides were characterized by predominately brief (0.2-2 sec) blocking events in comparison to the native toxin (20 sec). Histogram analysis revealed that mono-iodination of ChTX-Lq1 impairs blocking activity by adverse effects on both dissociation and association rate constants. Frequency density histograms of single channel blocking events provide a sensitive assay of toxin purity suitable for quantitating structure-activity relationships of charybotoxin derivatives.

Record Date Created: 1989/11/01 Record Date Completed: 1989/11/01

6776 (Item 6 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

086775595 PMID: 2473920 Charybotoxin blocks both Ca-activated K channels and Ca-independent K channels in rat brain synaptosomes.

Schneider M J; Rogowski R S; Krueger B K; Blaustein M P

Department of Physiology, University of Maryland School of Medicine, Baltimore 21201. FEBS Letters (NETHERLANDS) Jul 3 1989, 250 (2) p433-6, ISSN 0014-5793 Journal Code: 0155157

Contract/Grant No.: NS-16106; NS; NINDS; NS-16285; NS; NINDS; NS-20106; NS; NINDS; NS-20106. Record type: MEDLINE; Completed

Charybotoxin (ChTX), a 4.3 kDa polypeptide toxin from the venom of the scorpion *Leiurus quinquestratus*, blocks both a Ca-activated K channel (IC50 approximately 15 nM) and a Ca-independent voltage-gated K channel (IC50 approximately 40 nM) in rat brain synaptosomes. These results indicate that in this preparation ChTX is not specific for the Ca-activated K channel and suggest that there may be structural homology among the toxin-binding sites on various types of K channels.

Record Date Created: 1989/09/07 Record Date Completed: 1989/09/07

6779 (Item 9 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

110563434 PMID: 7513240 Charybotoxin and its effects on potassium channels.

Garcia M L; Knaus H G; Munujos P; Slaughter R S; Kaczorowski G J

Department of Membrane Biochemistry and Biophysics, Merck Research Laboratories, Rahway, New Jersey 07065, USA. American Journal of Physiology (UNITED STATES) Jul 1995, 269 (1 Pt 1) pC1-10, ISSN 0002-9513 Journal Code: 0370511 Publishing Model Print Document type: Journal Article; Review; Tutorial Languages: ENGLISH

Main Citation Owner: NLM Record type: MEDLINE; Completed

Over the last few years, a considerable amount of information has been obtained regarding K⁺ channels. Different areas of research have contributed to knowledge in this field. Charybotoxin (ChTX), a 37-amino acid peptide isolated from venom of the scorpion *Leiurus quinquestratus* var. hebraeus, represents a remarkable tool for studying K⁺ channels. With its use, it has been possible to purify the high-conductance Ca(2+)-activated K⁺ channel to homogeneity and determine the subunit composition of this channel. This has led to the discovery of an auxiliary beta-subunit that, when coexpressed with the pore-forming subunit, mSlo, alters the biophysical and pharmacological properties of this latter subunit. With the feasibility of producing large amounts of ChTX by recombinant techniques and the knowledge of the three-dimensional structure of the peptide, it has been possible to carry out site-directed mutagenesis studies and obtain a picture of the interaction surface of the toxin with two channels, maxi-K and Shaker, and to derive a picture of the complementary surface of the receptor in these two channels. Finally, ChTX, and the more selective K⁺ channel toxins that were subsequently discovered, have provided us with unique tools not only to determine the functional role that K⁺ channels play in target tissues but also to develop the molecular pharmacology of these channels. (76 Refs.)

Record Date Created: 1995/09/01 Record Date Completed: 1995/09/01

678111 (Item 11 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

08859199 PMID: 2482078 Charybotoxin is a new member of the K⁺ channel toxin family that includes dendrotoxin 1 and mast cell degranulating peptide.

Schweitz H; Bidard J N; Maes P; Lazdunski M

Centre de Biochimie, Centre National de la Recherche Scientifique, Université de Nice, France. Document type: Journal Article Languages: ENGLISH

Record type: MEDLINE; Completed

A polypeptide was identified in the venom of the scorpion *Leiurus quinquestratus* hebraeus by its potency to inhibit the high-affinity binding of the radiolabeled snake venom toxin dendrotoxin 1 (125I-DTX1) to its receptor site. It has been purified, and its properties investigated by different techniques were found to be similar to those of MCD and DTX1, two polypeptide toxins active on a voltage-dependent K⁺ channel. However, its amino acid sequence was determined, and it was shown that this toxin is in fact charybotoxin (ChTX), a toxin classically used as a specific tool to block one class of Ca2+-activated K⁺ channels. ChTX, DTX1, and MCD are potent convulsants and are highly toxic when injected.

intracerebroventricularly in mice. Their toxicities correlate well with their affinities for their receptors in rat brain. These three structurally different toxins release [³H]GABA from preloaded synaptosomes, the efficiency order being DTX1 greater than ChTX greater than MCD. Both binding and cross-linking experiments of ChTX to rat brain membranes and to the purified MCD/DTX1 binding protein have shown that the alpha-subunit (Mr = 76k-78k) of the MCD/DTX1-sensitive K⁺ channel protein also contains the ChTX binding sites. Binding sites for DTX1, MCD, and ChTX are in negative allosteric interaction. Our results show that charybotoxin belongs to the family of toxins which already includes the dendrotoxins and MCD, which are blockers of voltage-sensitive K⁺ channels. ChTX is clearly not selective for Ca2+-activated K⁺ channel.

Record Date Created: 1990/03/05 Record Date Completed: 1990/03/05

67718 (Item 18 from file: 5) DIALOG(R)File 5: Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

001134939 BIOSIS NO.: 19980143566

Consequence of the removal of evolutionary conserved disulfide bridges on the structure and function of charybotoxin and evidence that particular cysteine spacing governs specific disulfide bond formation

AUTHOR: Drakopoulou Eugenia; Vizzanova Jean; Neyton Jacques; Aniort Vincent ; Bouet Francios; Virelizier Henri; Menez Andre; Vila Claudio (Reprint)

AUTHOR ADDRESS: CEA, Dep. Ingénierie Etudes Protéines, Serv. Phys. Exp. Analyse, Saclay, 91190 Gif-sur-Yvette, France**; France

JOURNAL: Biochemistry 37 (5) p1292-1301 Feb. 3, 1998 1998 MEDIUM: print ISSN: 0006-2960 DOCUMENT TYPE: Article

RECORD TYPE: Abstract; LANGUAGE: English

ABSTRACT: Scorpion toxins are miniglobular proteins containing a common structural motif formed by an alpha-helix on one face, an antiparallel beta-sheet on the opposite face, and three disulfide bonds making up most of its internal volume. We have investigated the role of these evolutionary conserved bonds by replacing each couple of bridged cysteine residues of the scorpion charybotoxin by a pair of nonbridging L-alpha-aminobutyric acid (Aba) residues. Three analogues were obtained by solid-phase synthesis, Chab I, Chab II, and Chab III, containing the Aba residues in positions 7 and 28, 13 and 33, 17 and 35, respectively. Circular dichroism analysis showed that the purified Chab II acquired a conformation similar to that of charybotoxin, while the Chab I and Chab III possess decreased native-like characteristics. All analogues block single high-conductance Ca2+-activated K⁺ channels from rat skeletal muscle inserted into planar lipid bilayers, but with different potencies. Chab II is the most active analogue (KD = 8.0 X 10-8 M), with a 9-fold lower affinity as compared to native charybotoxin. Chab I and Chab III have, respectively, 180- and 380-fold lower affinity. Therefore, the removal of evolutionary conserved disulfide bridges does not prevent the toxin to adopt a functional and presumably native-like structure. However, removal of one disulfide bond affects the yields of formation of correct pairing between the remaining cysteine residues, and only Chab I preserves the ability to form the native disulfide pairings with high efficiency. This is the only analogue to preserve particular spacings of three and one residue between the cysteines, which have been described to be thermodynamically disfavor disulfide bond formation between the cysteines (Zhang, R. and Snyder, G. H. (1989) J. Biol. Chem. 264, 18472-18479). Therefore, we conclude that the position of the cysteine residues in the sequence of charybotoxin, by disfavoring specific pairings and favoring others, may govern selective formation of specific disulfide bonds, thus, explaining the efficient folding properties of Chab I and of native charybotoxin. The structural properties of the Chab analogues and the discovered role of the cysteine spacings have interesting implications in protein design and engineering.

67719 (Item 19 from file: 5) DIALOG(R)File 5: Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

000773556 BIOSIS NO: 19911911847

DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXIN A PEPTIDE BLOCKER OF POTASSIUM ION CHANNELS

AUTHOR: PARK C S; (Reprint); HAUSDORFF S F; MILLER C C

AUTHOR ADDRESS: HOWARD HUGHES MED INST GRADUATE DEP BIOCHEM, BRANDEIS UNIV, WALTHAM, MASS 02254, USA**; USA

JOURNAL: Proceedings of the National Academy of Sciences of the United States of America 88 (6) p2046-2050 1991 ISSN: 0027-8424 DOCUMENT TYPE: Article

ABSTRACT: A gene encoding charybotoxin (CTX), a K⁺ channel blocker from scorpion venom, was designed, synthesized, and expressed as a cleavable fusion protein in *Escherichia coli*. A sequence-specific protease, factor Xa, was used to cleave the fusion protein and thus release the toxin peptide. The recombinant toxin was purified, oxidized to form disulfide bonds, and treated to form N-terminal pyroglutamate. Recombinant CTX is identical to the native venom CTX with respect to high-performance liquid chromatography mobility, amino acid composition, and N-terminal modification. With single Ca2+-activated K⁺ channels as an assay system, recombinant CTX shows blocking and dissociation kinetics identical to the native venom toxin. The synthetic gene and high-level expression of functionally active CTX make it possible to study the fundamental mechanism of the toxin-ion channel interaction.

67722 (Item 22 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

08812992 PMID: 2531622 Effects of potassium channel toxins from *Leiurus quinquestratus hebraeus* venom on responses to cromakalim in rabbit blood vessels. Strong P N; Weir S W; Beech D J; Hiestand P; Kocher H P

1. The effects of fractionated *Lemurus quinquestratus* hebraeus venom on cromakalim-induced 86Rb+ efflux in rabbit aortic smooth muscle were examined. 2. Crude venom (0.1-10 micrograms ml-1) produced a concentration-dependent decrease of 1 microm cromakalim-induced 86Rb+ response. The maximum blocking activity attainable was approximately 60%. 3. Fractionation of crude venom by gel permeation chromatography and subsequent chromatography on a cation ion-exchange column, produced two fractions (X and XI), active in the 86Rb+-blocking assay. 4. Fraction XII contained charybdotoxin (approximately 85% pure). After a final high performance liquid chromatography (h.p.l.c.) purification step, the purified toxin failed to inhibit the cromakalim-stimulated 86Rb+ efflux although it was a potent inhibitor of A23187-induced K+ flux in human erythrocytes and the large conductance calcium-activated potassium channel in rabbit portal vein smooth muscle. 5. Subsequent purification of fraction X by h.p.l.c. yielded a minor peak which contained 86Rb+-blocking activity. This subfraction was also capable of inhibiting apamin-sensitive, angiotensin II-stimulated K+ flux in guinea-pig hepatocytes. 6. It is concluded that the potassium channel opened by cromakalim in rabbit aortic smooth muscle is not blocked by charybdotoxin but by another distinct toxin in the venom of *Lemurus quinquestratus* hebraeus.

Record Date Created: 19900125 Record Date Completed: 19900125

67729 (Item 29 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog . All rts. reserv.

07773735 PMID: 2433153

Identification of two toxins from scorpion (*Lemurus quinquestratus*) venom which block distinct classes of calcium-activated potassium channel.

Castle N; Strong P N

FEBS Letters (NETHERLANDS) Dec 1 1986, 209 (1) p117-21; ISSN 0014-5793 Journal Code: 0155157

Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM

Record type: MEDLINE; Completed

Two polypeptide toxins from scorpion (*Lemurus quinquestratus*) venom which block distinct classes of calcium-activated potassium channels have been identified and partially purified. One toxin, at 50-100 ng/ml, blocks apamin-sensitive potassium fluxes in hepatocytes and inhibits [125I]monooctapeptide binding. The other, more basic, toxin blocks apamin-insensitive potassium fluxes in erythrocytes at 200 ng/ml and, to our knowledge, is the first toxin shown to block the erythrocyte calcium-activated potassium channel with high affinity. The possible co-identity of this latter toxin with charybdotoxin is discussed.

Record Date Created: 19870220 Record Date Completed: 19870220

67733 (Item 33 from file: 5) DIALOG(R)File 5: Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0006395608 BIOSIS NO.: 198936104699

LEUROUS-QUINQUESTRATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES

AUTHOR: SORPSEN R G (Reprint); SCHNEIDER M J; BLAUSTEIN M P

AUTHOR ADDRESS: MED DEP, DIV ENVIRONMED AND TOXICOL, JEFFERSON MED COLL, PHILADELPHIA, PA 19107, USA**USA

JOURNAL: Biophysical Journal 55 (2 PART 2); p560A 1989 CONFERENCE/MEETING: THIRTY-THIRD ANNUAL MEETING OF THE BIOPHYSICAL SOCIETY, CINCINNATI, OHIO, USA, FEBRUARY 12-16, 1989. BIOPHYS J ISSN: 0006-3495

DOCUMENT TYPE: Meeting RECORD TYPE: Citation LANGUAGE: ENGLISH

10895806 PMID: 7533951

67736 (Item 36 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog . All rts. reserv.

Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Lemurus quinquestratus* hebraeus.

Marshall D L; Vatrapour H; Harvey A L; Boyot P; Pinkasfeld S; Doljansky Y; Bouet F; Menet A

Department of Physiology and Pharmacology, University of Strathclyde, Glasgow, U.K.

Toxincon - official journal of the International Society on Toxicology (ENGLAND) Nov 1994, 32 (11) p1433-43; ISSN 0041-0101 Journal Code: 1307333 Publishing Model Print . Document type: Journal Article Languages: ENGLISH

Main Citation Owner: NLM Record type: MEDLINE; Completed

The scorpion venom *Lemurus quinquestratus* hebraeus was fractionated by chromatography in order to isolate toxins that affected binding of radiolabelled dendrotoxin to K+ channel proteins on synaptosomal membranes and that facilitated acetylcholine release in chick biventer cervicus nerve-muscle preparations. In addition to the previously characterized charybdotoxin, three toxins were isolated: 14.2, 15.1 and 18.2. Toxin 14.2 has a blocked N-terminus and because of low quantities, it has not been sequenced; 15.1 is a newly sequenced toxin of 36 residues with some overall homology to charybdotoxin and noxiustoxin; 18.2 is identical to charybdotoxin-2. The apparent Ki against dendrotoxin binding were: charybdotoxin - 3.8 nm, 14.2, 150 nm; 15.1, 50 nm, and 18.2, 0.25 nm. Toxin 14.2 (75 nm-1.5 microm) had a presynaptic

facilitatory effect on neuromuscular preparations. Toxin 15.1 augmented responses to direct muscle stimulation, probably because it blocked Ca(2+)-activated K+ currents in muscle fibres. Toxin 18.2 (charybdotoxin-2) had a potent presynaptic facilitatory action, with less effect on direct muscle stimulation. This contrasts with the relatively weak neuromuscular effects of the highly homologous charybdotoxin. On a Ca(2+)-activated K+ current in mouse motor nerve endings, charybdotoxin and toxin 18.2 produced maximal block at around 100 nM, whereas 15.1 was inactive at 300 nM. Charybdotoxin can increase quantal content, but this is more likely to result from block of voltage-dependent K+ channels than Ca(2+)-activated channels; the increase in transmitter release occurred in conditions in which little IKCa would be present; higher concentration of charybdotoxin and longer exposure times were required to increase transmitter release than those needed to block IKCa, and the facilitatory effects of charybdotoxin and toxin 18.2 correlated more with their effects on dendrotoxin binding than on block of IKCa.

Record Date Created: 19900410 Record Date Completed: 19950410

67745 (Item 45 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog . All rts. reserv.

08257702 PMID: 2453055

Purification, sequence, and model structure of charybdotoxin, a potent selective inhibitor of calcium-activated potassium channels.

Gimenez-Gallego G; Navia M A; Reuben J P; Katz G M; Kaczorowski G J; Garcia M L

Department of Growth Factor Research, Merck Sharp & Dohme Research Laboratories, Rahway, NJ 07065

Proceedings of the National Academy of Sciences of the United States of America (UNITED STATES) May 1988, 85 (10) p329-33; ISSN 0027-8424 Journal Code: 7505876 Publishing Model Print Document type: Journal Article

Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed

Charybdotoxin (ChTX), a protein present in the venom of the scorpion *Leiurus quinquestratus* var. hebraeus, has been purified to homogeneity by a combination of ion-exchange and reversed-phase chromatography. Polycrylamide gel electrophoresis, amino acid analysis, and complete amino acid sequence determination of the pure protein reveal that it consists of a single polypeptide chain of 4.3 kDa. Purified ChTX is a potent and selective inhibitor of the approximately 220-pS Ca2+-activated K+ channel present in GH3 anterior pituitary cells and primary bovine aortic smooth muscle cells. The toxin reversibly blocks channel activity by interacting at the external pore of the channel protein with an apparent Kd of 2.1 nM. The primary structure of ChTX is similar to a number of neurotoxins of diverse origin, which suggests that ChTX is a member of a superfamily of proteins that modify ion-channel activities. On the basis of this similarity, the three-dimensional structure of ChTX has been modeled from the known crystal structure of alpha-bungarotoxin. These studies indicate that ChTX is useful as a probe of Ca2+-activated K+-channel function and suggest that the proposed tertiary structure of ChTX may provide insight into the mechanism of channel block.

Record Date Created: 19880622 Record Date Completed: 19980622

67752 (Item 52 from file: 5) DIALOG(R)File 5: Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0010206283 BIOSIS NO.: 199698674126

Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin (Reprint)

AUTHOR: Dohle Timothy R; Duggan Brendan M; Pennington Michael W; Byrnes Michael E; Ken William R; Norton Raymond S (Reprint)

AUTHOR ADDRESS: NMR Lab, Biomolecular Research Inst, 381 Royal Parade, Parkville, VIC 3052, Australia**Australia

JOURNAL: Biophysical Acta 1292 (1): p31-38 1996 1996 ISSN: 0006-3002 DOCUMENT TYPE: Article

RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: Charybdotoxin is a 37-residue polypeptide toxin from scorpion venom, which acts by blocking voltage-gated and Ca2+-activated K+ channels. We have synthesized charybdotoxin and three mono-substituted analogues using an Fmoc-Ibu protocol. The Phe2 Iwdaw Tyr analogue was chosen to introduce a site for Tyr iodination which was distinct from the K+ channel binding surface, while the Glu-12 Iwdaw Gln and Arg-19 Iwdaw His analogues were studied to probe the roles of charged residues at these positions in the structure and activity of the toxin. The synthetic native molecule was equipotent with natural toxin in inhibiting the human erythrocyte Ca2+-dependent K+ channel. The affinities of all three analogues for the erythrocyte K+ channel were, slightly reduced with the Arg-19 Iwdaw His analogue showing the greatest increase in IC-50 (2.3-fold). Two-dimensional 1H-NMR studies of these analogues showed that all three had structures very similar to those of the native molecule. The most significant perturbation was associated with the Glu-12 to Gln substitution, which appeared to destabilise the N-terminal half of the alpha-helix, possibly due to the weakening of an N-terminal helix capping interaction which is apparent from our NMR data. His-21 has a pKa more than one unit below the value for a non-interacting histidine. Possible reasons for this are that the imidazolium side chain is partly buried and is located near positively charged moieties. Thus, His-21 would be neutral at physiological pH, where charybdotoxin binds to the potassium channel.

67762 (Item 62 from file: 5) DIALOG(R)File 5: Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0007742476 BIOSIS NO.: 199119125367

THREE-DIMENSIONAL STRUCTURE OF NATURAL CHARYBDOTOXIN IN AQUEOUS SOLUTION BY PROTON NMR CHARYBDOTOXIN POSSESSES A STRUCTURAL MOTIF FOUND IN OTHER SCORPION TOXINS

AUTHOR: BONTEMPS F (Reprint); ROUMESTAND C; BOYOT P; GILQUIN B; DOLJANSKY Y; MENEZA; TOMAF

AUTHOR ADDRESS: SERV DE BIOCHIM DES PROTEINES, LAB D'INGENIERIE DES PROTEINES BAT 152, CEN-SACLAY, F-91191 GIF-SUR-YVETTE, FRANCE

JOURNAL: European Journal of Biochemistry 156 (1): p19-28 1991 ISSN: 0014-2956 DOCUMENT TYPE: Article

RECORD TYPE: Abstract LANGUAGE: ENGLISH

ABSTRACT: A 600-MHz proton NMR study of natural charybotoxin, a toxin acting on K-channels, is reported. The analysis of NOEs and of backbone coupling constants showed the existence of an alpha-helix (residues 10-19) and of an antiparallel beta-sheet in the 26-35 part. Three-dimensional structures were generated by distance geometry, using a set of 114 inter-residual calibrated constraints (63 sequential, 47 medium and long range, 4 hydrogen bonds) and 29 PHI angles. These structures show that charybotoxin is composed of a beta-sheet linked to an alpha-helix by two disulphide bridges and to an extended fragment by the third disulphide bridge. Comparison with the other known structures of long and short scorpion toxins shows that this structural motif is common to all these proteins.

9/6/1 (Item 1 from file: 5) 0007115178 BIOSIS NO.: 199089033069 ANALYSIS OF THE BLOCKING ACTIVITY OF CHARYBDOTOXIN HOMOLOGS AND IODINATED DERIVATIVES AGAINST CALCIUM-ACTIVATED POTASSIUM CHANNELS 1989

9/6/2 (Item 2 from file: 155) 08751830 PMID: 2475748

Analysis of the blocking activity of charybotoxin homologs and iodinated derivatives against Ca^{2+} -activated K+ channels. Aug 1989

9/6/3 (Item 3 from file: 155) 08675535 PMID: 2473920

Charybotoxin is a new member of the K+ channel toxin family that includes dendrotoxin 1 and mast cell degranulating peptide. Dec 12 1989

9/6/4 (Item 4 from file: 155) 088559199 PMID: 2482078

Charybotoxin is a new member of the K+ channel toxin family that includes dendrotoxin 1 and mast cell degranulating peptide. Dec 12 1989

9/6/5 (Item 5 from file: 5) 0007133033 BIOSIS NO.: 199089050924

CHARYBDOTOXIN IS A NEW MEMBER OF THE POTASSIUM CHANNEL TOXIN FAMILY THAT INCLUDES DENDROTOXIN 1 AND MAST CELL DEGRANULATING PEPTIDE 1989

9/6/6 (Item 6 from file: 5) 0006941512 BIOSIS NO.: 199038119403

CHARYBDOTOXIN IS A NEW MEMBER OF THE TOXIN FAMILY THAT INCLUDES DENDROTOXIN 1 AND MCD AND BLOCKS DENDROTOXIN-SENSITIVE VOLTAGE ACTIVATED POTASSIUM CHANNELS 1990

9/6/7 (Item 7 from file: 5) 0011349339 BIOSIS NO.: 199800143586

Consequence of the removal of evolutionarily conserved disulfide bridges on the structure and function of charybotoxin and evidence that particular cysteine spacing govern specific disulfide bond formation 1998

9/6/8 (Item 8 from file: 155) 08571405 PMID: 2468931

Effect of some potassium channel blockers on contractile responses of the rabbit aorta. Feb 1989

9/6/9 (Item 9 from file: 5) 0006395808 BIOSIS NO.: 198936104699

LEIURUS-QUINQUESTRATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES 1989

9/6/10 (Item 10 from file: 155) 10823421 PMID: 7819188

NMR sequential assignments and solution structure of charybotoxin, a small scorpion toxin that blocks chloride channels. Jan 10 1995

9/6/11 (Item 11 from file: 5) 0009648272 BIOSIS NO.: 199598116660

NMR Sequential Assignments and Solution Structures of Charybotoxin, a Small Scorpion Toxin That Blocks Chloride Channels 1995

9/6/12 (Item 12 from file: 155) 10895806 PMID: 7533951

Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestratus hebreus*. Nov 1994

9/6/13 (Item 13 from file: 5) 0009575215 BIOSIS NO.: 198598043048

Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestratus hebreus* 1994

9/6/14 (Item 14 from file: 5) 0008709107 BIOSIS NO.: 199395011373

Noxitoxin and leuoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels 1992

9/6/15 (Item 15 from file: 5) 0010206293 BIOSIS NO.: 19968674126

Synthesis and structural characterisation of analogues of the potassium channel blocker charybotoxin 1996

9/6/16 (Item 16 from file: 5) 0008934542 BIOSIS NO.: 19949705327

Synthesis of charybotoxin and of two N-terminal truncated analogues : Structural and functional characterization 1993

9/6/17 (Item 17 from file: 5) 0010206293 BIOSIS NO.: 19968674126

Synthesis and structural characterisation of analogues of the potassium channel blocker charybotoxin 1996

9/6/18 (Item 18 from file: 5) 0009025088 BIOSIS NO.: 199497046373 Toxin pharmacology of the large-conductance Ca^{2+} -activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture 1993

12/6/1 (Item 1 from file: 155) 11322494 PMID: 8645186

A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator*. May 1 1996

13/6/1 (Item 1 from file: 155) 11322494 PMID: 8645186

A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator*. May 1 1996

13/6/2 (Item 1 from file: 5) 0010380507 BIOSIS NO.: 1996869014567

A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator* 1996

16/6/1 (Item 1 from file: 155) 12140757 PMID: 9438859

Solution structure and proposed binding mechanism of a novel potassium channel toxin kappa-conotoxin PVIIA. Dec 15 1997

16/6/2 (Item 1 from file: 5) 0011293829 BIOSIS NO.: 199800088076

Solution structure and proposed binding mechanism of a novel potassium channel toxin kappa-conotoxin PVIIA 1997

19/6/1 (Item 1 from file: 155) 10204659 PMID: 76856355

An activator of calcium-dependent potassium channels isolated from a medicinal herb. Jun 22 1993

19/6/2 (Item 2 from file: 155) 15271073 PMID: 15051409

Antigenic polymorphism of the "star" scorpion toxins able to block K+ channels. Mar 15 2004

19/6/3 (Item 3 from file: 5) 0009645783 BIOSIS NO.: 198783124674

BLOCKING AGENTS OF CALCIUM-ACTIVATED POTASSIUM CHANNELS IN CULTURED MEDULLARY THICK ASCENDING LIMB CELLS 1987

19/6/4 (Item 4 from file: 155) 07821071 PMID: 2435161

Blocking agents of Ca^{2+} -activated K+ channels in cultured medullary thick ascending limb cells. Feb 1 1987

19/6/5 (Item 5 from file: 155) 17947714 PMID: 15695820

BmP09, a "big chain" scorpion peptide blocker of BK channels. Apr 15 2005

19/6/6 (Item 6 from file: 5) 0008848545 BIOSIS NO.: 1993936012961

Calcium-activated potassium transport in erythrocytes: Comparison of binding and transport inhibition by scorpion toxins 1993

19/6/7 (Item 7 from file: 155) 10517036 PMID: 8297371

Chemical synthesis and structure-function studies of margatoxin, a potent inhibitor of voltage-dependent potassium channel in human T lymphocytes. Jan 28 1994

19/6/8 (Item 8 from file: 155) 08816237 PMID: 2483347

Characterization of high affinity binding sites for charybotoxin in sarcoplasmic membranes from bovine aortic smooth muscle. Evidence for a direct association with the high conductance calcium-activated potassium channel. Dec 15 1989

19/6/9 (Item 9 from file: 155) 11411894 PMID: 8705835

Characterization of a new peptide from *Tityus serrulatus* scorpion venom which is a ligand of the apamin-binding site. Jul 15 1996

19/6/10 (Item 10 from file: 155) 16510736 PMID: 15189853

Computational simulations of interactions of scorpion toxins with the voltage-gated potassium ion channel. Jun 2004

19/6/11 (Item 11 from file: 5) 0011960511 BIOSIS NO.: 200400331297

Computational simulations of interactions of scorpion toxins with the voltage-gated potassium ion channel 2004

19/6/12 (Item 12 from file: 155) 12172929 PMID: 9477955

Consequence of the removal of evolutionary conserved disulfide bridges on the structure and function of charybotoxin and evidence that

particular cysteine spacings govern specific disulfide bond formation. Feb 3 1998

19/6/13 (Item 13 from file: 155) 110191649 PMID: 7545007

Cross-linking of charybotoxin to high-conductance calcium-activated potassium channels: identification of the covalently modified toxin residue. Aug 28 1995

19/6/14 (Item 14 from file: 155) 11196506 PMID: 8531201

Ca(2+)-activated K+ channels of human and rabbit erythrocytes display distinctive patterns of inhibition by venom peptide toxins. Sep 1995

19/6/15 (Item 15 from file: 155) 10145953 PMID: 7682555

Ca(2+)-activated K+ transport in erythrocytes. Comparison of binding and transport inhibition by scorpion toxins. Apr 25 1993

19/6/16 (Item 16 from file: 155) 09358728 PMID: 1706481

196/60 (Item 60 from file: 5) 0014975010 BIOSIS NO.: 200400345799
Solution structure of BmBKT₁, a new K₊-channel blocker from the Chinese scorpion *Buthus martensi* Karsch 2004

196/61 (Item 61 from file: 155) 14028726 PMID: 11790849
Solution structure of a K₊-channel blocker from the scorpion *Tityus cambridgei*. Feb 2002

196/62 (Item 62 from file: 155) 11808576 PMID: 962103
Solution structure for Pandinus toxin K-alpha (PFTX-K alpha), a selective blocker of A-type potassium channels. Mar 11 1997

196/63 (Item 63 from file: 155) 12075374 PMID: 965990
Solution structure of Tskapa, a charybdotoxin-like scorpion toxin from *Tityus serratus* with high affinity for apamin-sensitive Ca₂₊-activated K₊ channels. Nov 1997

196/64 (Item 64 from file: 5) 0010328748 BIOSIS NO.: 199608796381
Solution structure of SHK toxin, a novel potassium channel inhibitor from a sea anemone 1996

196/65 (Item 65 from file: 155) 11968450 PMID: 9252467
Sapecin B, a novel ly toxin, blocks macroscopic K₊ currents in the GH3 rat pituitary cell line. Jul 1997

196/66 (Item 66 from file: 5) 0011067804 BIOSIS NO.: 199799701864
Sapecin B, a novel ly toxin, blocks macroscopic K₊ currents in the GH3 rat pituitary cell line 1997

196/67 (Item 67 from file: 5) 0012839056 BIOSIS NO.: 200100010905
Structure determinants of scorpion toxin affinity: The charybdotoxin (alpha-KTX) family of K₊-channel blocking peptides BOOK TITLE: Reviews of Physiology Biochemistry and Pharmacology 2000

196/68 (Item 68 from file: 155) 12909571 PMID: 10857399
Structural determinants of scorpion toxin affinity: the charybdotoxin (alpha-KTX) family of K₊-channel blocking peptides. 2000

196/69 (Item 69 from file: 155) 12762762 PMID: 10707030
Structural and functional differences of two toxins from the scorpion *Pandinus imperator*. Mar 1 2000

196/70 (Item 70 from file: 5) 0012447874 BIOSIS NO.: 200000016187
Structural and functional differences of two toxins from the scorpion *Pandinus imperator* 2000

196/71 (Item 71 from file: 5) 0010968956 BIOSIS NO.: 19979623016
Structural analysis of a two disulfide bridge analogue of a scorpion toxin 1997

196/72 (Item 72 from file: 155) 10377674 PMID: 8253752
Synthesis and characterization of Kalotoxin. Is the 26-32 sequence essential for potassium channel recognition? Dec 15 1993

196/73 (Item 73 from file: 155) 10341798 PMID: 7693459
Synthesis of charybdotoxin and of two N-terminal truncated analogues. Structural and functional characterisation. Oct 1 1993

196/74 (Item 74 from file: 155) 11245554 PMID: 8547346
Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin. Jan 4 1996

196/75 (Item 75 from file: 155) 11180566 PMID: 7576559
Topology of the pore-region of a K₊-channel revealed by the NMR-derived structures of scorpion toxins. Nov 1995

196/76 (Item 76 from file: 155) 10582194 PMID: 7514038
Tremorgenic indole alkaloids potently inhibit smooth muscle high-conductance calcium-activated potassium channels. May 17 1994

196/77 (Item 77 from file: 155) 14501863 PMID: 12445473
Two novel toxins from the Amazonian scorpion *Tityus cambridgei* that block Kv1.3 and Shaker B K₊-channels with distinctly different affinities. Dec 16 2002

196/78 (Item 78 from file: 155) 12343405 PMID: 8655636
Two similar peptides from the venom of the scorpion *Pandinus imperator*, one highly effective blocker and the other inactive on K₊-channels. May 1998

196/79 (Item 79 from file: 5) 0011532158 BIOSIS NO.: 199800326405
Two similar peptides from the venom of the scorpion *Pandinus imperator*, one highly effective blocker and the other inactive on K₊-channels 1998

196/80 (Item 80 from file: 155) 11603837 PMID: 8913348
Three new toxins from the scorpion *Pandinus imperator* selectively block certain voltage-gated K₊-channels. Nov 1996

196/81 (Item 81 from file: 5) 0010694277 BIOSIS NO.: 199799328337
Three new toxins from the scorpion *Pandinus imperator* selectively block certain voltage-gated K₊-channels 1996

24/6/1 (Item 1 from file: 155) 11928755 PMID: 9208943
Anti-insect toxin 5 (AaIT5) from *Androctonus australis*. Jun 1 1997

24/6/2 (Item 2 from file: 5) 001097935 BIOSIS NO.: 199799611995
Anti-insect toxin 5 (AaIT5) from *Androctonus australis* 1997

24/6/3 (Item 3 from file: 357) 0206373 DBR Accession No.: 97-01494
Control of insects, acar and nematodes - recombinant *Autographa californica* nuclear-polyhedrosis virus vector-mediated scorpion, snail, mite or spider venom toxin gene expression for use as a biological control agent 1996

24/6/4 (Item 4 from file: 155) 11146500 PMID: 8533143
Positive cooperativity among insecticidal scorpion neurotoxins. Aug 1995

24/6/5 (Item 5 from file: 5) 0009989127 BIOSIS NO.: 199598456360
Positive cooperativity among insecticidal scorpion neurotoxins 1995

24/6/6 (Item 6 from file: 155) 12339622 PMID: 9652392
A depressant insect-selective toxin anabg from the venom of the scorpion structure/function characterization. May 15 1998

24/6/7 (Item 7 from file: 5) 0011517804 BIOSIS NO.: 199800312051
A depressant insect-selective toxin anabg from the venom of the scorpion Leitus quinquestriatus hebraeus; Purification and structure/function characterization 1998

24/6/8 (Item 8 from file: 155) 10940776 PMID: 7722081
Direct effects of recombinant nuclear polyhedrosis viruses on selected nontarget organisms. Apr 1995

24/6/9 (Item 9 from file: 5) 0009776161 BIOSIS NO.: 199598244443
Direct effects of recombinant nuclear polyhedrosis viruses on selected nontarget organisms 1995

24/6/10 (Item 10 from file: 5) 0011296947 BIOSIS NO.: 2003025566
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nuclear-polyhedrovirus expressing the scorpion toxin Lqh112. 2003

24/6/11 (Item 11 from file: 357) 0313302 DBR Accession No.: 2003-14442
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nuclear-polyhedrovirus expressing the scorpion toxin Lqh112 - scorpion venom toxin expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

24/6/12 (Item 12 from file: 155) 12084031 PMID: 9374355
Interactions of recombinant and wild-type baculoviruses with classical insecticides and pyrethroid-resistant tobacco budworm (Lepidoptera: Noctuidae). Oct 1997

24/6/13 (Item 13 from file: 5) 0009921314 BIOSIS NO.: 199598389147
Insect sodium channel as the target for insect-selective neurotoxins from scorpion venom BOOK TITLE: ACS Symposium Series, Molecular action of insecticides on ion channels 1995

24/6/14 (Item 14 from file: 5) 0009184111 BIOSIS NO.: 199497205396
The insect sodium channel as the target for the insect selective neurotoxins from scorpion venom 1994

24/6/15 (Item 15 from file: 5) 0010929668 BIOSIS NO.: 199799563728
Insect tolerance to a neurotoxic polypeptide: Pharmacokinetic and pharmacodynamic aspects 1997

24/6/16 (Item 16 from file: 155) 14493200 PMID: 11782289
Isolation and characterization of a novel lepidopteran-selective toxin from the venom of South Indian red scorpion, *Mesobuthus tamulus*. 2001

24/6/17 (Item 17 from file: 5) 0013739895 BIOSIS NO.: 200200387406
Isolation and characterization of a novel lepidopteran-selective toxin from the venom of South Indian red scorpion, *Mesobuthus tamulus* 2001

24/6/18 (Item 18 from file: 357) 0264696 DBR Accession No.: 2001-04723
New polynucleotides encoding scorpion venom potassium channel-agonist proteins for producing e.g. of insect-tolerant transgenic plants for controlling insect pest damage and parasitic worm infections - scorpion venom potassium channel-agonist protein genes useful for constructing transgenic plant with insect resistance 2000

24/6/19 (Item 19 from file: 357) 0264374 DBR Accession No.: 2001-04128
New isolated polynucleotide encoding a scorpion toxin for treating epilepsy, degenerative disorders such as Huntington's disease, and neuronal death following stroke, and for creating plants that are insect tolerant - transgenic plant construction with insect resistance and gene therapy 2000

24/6/20 (Item 20 from file: 5) 00086164579 BIOSIS NO.: 19934509567
Potential of baculoviruses expressing a scorpion toxin and an esterase in agriculture 1993

24/6/21 (Item 21 from file: 357) 0153538 DBR Accession No.: 93-11590

Potential of baculoviruses expressing a scorpion toxin and an esterase in agriculture - use of recombinant baculovirus with a *Heliothis virescens* juvenile-hormone-esterase as an insect biological control agent (conference abstract) 1993

24/6/22 (Item 22 from file: 155) 12317587 PMID: 9627406

Rapid purification and molecular modeling of *AaIT* peptides from venom of *Androctonus australis*. 1998

24/6/23 (Item 23 from file: 5) 0011525675 BIOSIS NO.: 19980031922

Rapid purification and molecular modeling of *AaIT* peptides from venom of *Androctonus australis* 1998

24/6/24 (Item 24 from file: 5) 0014912774 BIOSIS NO.: 200400283531

Scorpion toxins 2004

24/6/25 (Item 25 from file: 5) 0007297260 BIOSIS NO.: 199090081739

A SCORPION VENOM NEUROTOXIN PARALYTIC TO INSECTS THAT AFFECTS SODIUM CURRENT INACTIVATION PURIFICATION PRIMARY STRUCTURE AND MODE OF ACTION 1990

24/6/26 (Item 26 from file: 155) 09075706 PMID: 2383565

A scorpion venom neurotoxin paralytic to insects that affects sodium current inactivation: purification, primary structure, and mode of action. Jun 26 1990

24/6/27 (Item 27 from file: 5) 0007645125 BIOSIS NO.: 199191028016

THE TOLERANCE OF LEPIDOPTEROUS LARVAE TO AN INSECT SELECTIVE NEUROTOXIN 1990

24/6/28 (Item 28 from file: 5) 0009069061 BIOSIS NO.: 199497090346

Variability among insect sodium channels revealed by selective neurotoxins 1994

24/6/29 (Item 29 from file: 5) 0010988996 BIOSIS NO.: 199799623056

Two novel short insectotoxins from the Asian scorpions *Butus martensi* and *Butus tamulus* 1997

24/7/6 (Item 6 from file: 155) DIALOG(R)File 155: MEDLINE(R)(c) format only 2005 Dialog. All its. reserv.

12336622 PMID: 9652392

A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus--purification and structure/function characterization. Moskowitz H; Hermann R; Jones A D; Hammock B D

Department of Entomology, University of California, Davis 95616, USA. European journal of biochemistry / FEBS (GERMANY) May 15, 1998, 254, (1) p44-9, ISSN 0014-2956 Journal Code: 0107600 Contract/Grant No.: P 30 ES05707; ES; NIEHS Publishing Model Print Document type: Journal Article

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The scorpion Lqh1T5-derived excitatory and depressant insect-selective polypeptide neurotoxins modify sodium conductance in insect neuronal membranes and differ greatly in their primary structures and symptoms induced in blow fly larvae. We report here the purification and characterization of a new insect selective toxin Lqh1T5. Lqh1T5 is more similar to the excitatory toxins in its mode of action and the depressant toxins in its primary structure. This toxin is a single polypeptide composed of 61 amino acids that are cross linked by four disulfide bonds. When Lqh1T5 is injected into blow fly larvae, a fast contraction paralysis occurs without depressant activity. No mammalian toxicity was detected by subcutaneous or intracranial injections of this toxin into mice. Sequence comparison of Lqh1T5 and known depressant toxins shows a high degree of similarity among the amino acids located on the C-terminus of the toxins. However, there are some clear differences in the amino acids located close to the N-terminus of the toxins. By the aid of homology modeling, we demonstrated that these amino acids have the same orientation in the tertiary structure of the molecule and are exposed to the environment. The change in the mode of action of Lqh1T5 (no depressant activity) by substitutions of a few amino acids located on a specific exposed area of the toxin shed a new light on the structure/function relationship of scorpion toxins. These results caution that similarity in the mechanism of action of scorpion toxins does not always follow from an overall similarity in sequence. Record Date Created: 19980728 Record Date Completed: 19980728

26/6/1 (Item 1 from file: 155) 11928755 PMID: 9208943

Anti-insect toxin 5 (AaIT5) from *Androctonus australis*. Jun 1 1997

26/6/2 (Item 2 from file: 5) 0010977935 BIOSIS NO.: 199799611985

Anti-insect toxin 5 (AaIT5) from *Androctonus australis*. 1997

26/6/3 (Item 3 from file: 155) 12336622 PMID: 9652392

A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus--purification and structure/function characterization. May 15 1998

26/6/4 (Item 4 from file: 5) 0011517804 BIOSIS NO.: 199800312051

A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus. Purification and structure/function characterization 1998

26/6/5 (Item 5 from file: 5) 0014296947 BIOSIS NO.: 200300255666 Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2. 2003

26/6/6 (Item 6 from file: 357) 0313302 DBR Accession No.: 2003-14442

Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2 - scorpion venom toxin expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

26/6/7 (Item 7 from file: 5) 0007297260 BIOSIS NO.: 199090081739

A SCORPION VENOM NEUROTOXIN PARALYTIC TO INSECTS THAT AFFECTS SODIUM CURRENT INACTIVATION PURIFICATION PRIMARY STRUCTURE AND MODE OF ACTION 1990

26/6/8 (Item 8 from file: 155) 09075706 PMID: 2383565

A scorpion venom neurotoxin paralytic to insects that affects sodium current inactivation: purification, primary structure, and mode of action. Jun 26 1990

26/6/9 (Item 9 from file: 5) 0009059061 BIOSIS NO.: 199497090346

Variability among insect sodium channels revealed by selective neurotoxins 1994

28/6/1 (Item 1 from file: 357) 0313302 DBR Accession No.: 2003-14442

Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2 - scorpion venom toxin expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

29/6/1 (Item 1 from file: 5) 0014419169 BIOSIS NO.: 200300377888

Scorpion toxins 2003

29/6/2 (Item 1 from file: 357) 0313302 DBR Accession No.: 2003-14442

Effect of signal sequences and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2 - scorpion venom toxin expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

29/6/3 (Item 2 from file: 357) 0256837 DBR Accession No.: 2000-11327

New nucleic acid fragment encoding a scorpion toxin that is potassium channel-agonist, useful for creating transgenic plants that are more insect-tolerant - method is useful for producing transgenic plant with insect resistance 2000

33/6/1 (Item 1 from file: 5) 0014296947 BIOSIS NO.: 200300255666

Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2. 2003

33/6/2 (Item 1 from file: 357) 0313302 DBR Accession No.: 2003-14442

Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2 - scorpion venom toxin expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

34/6/1 (Item 1 from file: 5) 0014419169 BIOSIS NO.: 200300377888

Scorpion toxins 2003

34/6/2 (Item 2 from file: 5) 0014296947 BIOSIS NO.: 200300255666

Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqh1T2. 2003

34/6/3 (Item 2 from file: 357) 0256837 DBR Accession No.: 2000-11327

New nucleic acid fragment encoding a scorpion toxin that is potassium channel-agonist, useful for creating transgenic plants that are more insect-tolerant - method is useful for producing transgenic plant with insect resistance 2000

43/6/1 (Item 1 from file: 155) 09694018 PMID: 1796479

The cDNA sequence of a depressant insect selective neurotoxin from the scorpion *Butthotus judaicus*. 1991

43/6/2 (Item 2 from file: 5) 00168178 BIOSIS NO.: 199293011769

THE cDNA SEQUENCE OF A DEPRESSANT INSECT SELECTIVE NEUROTOXIN FROM THE SCORPION BUTTHOTUS-JUDAICUS 1991

43/6/3 (Item 3 from file: 155) 12780322 PMID: 10708793

Cloning and characterization of a cDNA sequence encoding the precursor of a chirotoxin-like peptide from the Chinese scorpion *Butthotus martensi* Karsch. Aug 2000

43/6/4 (Item 4 from file: 5) 0012519314 BIOSIS NO.: 200000237627

Cloning and characterization of a cDNA sequence encoding the precursor of a chirotoxin-like peptide from the Chinese scorpion *Butthotus martensi* Karsch 2000

43665 (Item 5 from file: 5) 0014778747 BIOSIS NO.: 200400145408
Cytotoxic and apoptotic effects of scorpion *Leurus quinquestriatus* venom on 293T and C2C12 eukaryotic cell lines. 2003

43666 (Item 6 from file: 15) 10074904 PMID: 14431601
Depressant insect selective neurotoxins from scorpion venom: chemistry, action, and gene cloning. 1993

43667 (Item 7 from file: 5) 0011914255 BIOSIS NO.: 199900174585
Dynamic diversification from a putative common ancestor of scorpion toxins affecting sodium, potassium, and chloride channels 1989

43668 (Item 8 from file: 5) 0007735566 BIOSIS NO.: 199191118447
DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXINA PEPTIDE BLOCKER OF POTASSIUM CHANNELS 1991

43669 (Item 9 from file: 5) 12095528 PMID: 9395089
Influence of a NH2-terminal extension on the activity of KTx2, a K⁺ channel blocker purified from *Androctonus australis* scorpion venom. Nov 3 1997

43670 (Item 10 from file: 5) 0014296947 BIOSIS NO.: 20030255666
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqH72. 2003

43671 (Item 11 from file: 357) 0313302 DBR Accession No.: 2003-14442
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqH72 - scorpion venom toxin expression in nuclear polyhedrosis virus for biological control agent strain improvement 2003

43672 (Item 12 from file: 155) 12401430 PMID: 9714546
Evidence for a new class of scorpion toxins active against K⁺-channels. Jul 24 1998

43673 (Item 13 from file: 155) 11211987 PMID: 7498537
Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae. Dec 4 1995

43674 (Item 14 from file: 5) 0010170013 BIOSIS NO.: 199699637846
Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae 1995

43675 (Item 15 from file: 357) 01914143 DBR Accession No.: 96-02205
Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae - *Autographa californica* nucleopolyhedrosis virus vector allows insecticide expression in insect cell culture or larva 1995

43676 (Item 16 from file: 5) 0012082499 BIOSIS NO.: 199800342159
Genomic organization of three neurotoxins active no small conductance Ca²⁺-activated potassium channels from the scorpion *Buthus martensi Karsch* 1989

43677 (Item 17 from file: 155) 18326469 PMID: 15966742
Genetic polymorphism and expression of a highly potent scorpion depressant toxin enable refinement of the effects on insect Na channels and illuminate the key role of Asn-58. Jun 28 2005

43678 (Item 18 from file: 5) 0015265895 BIOSIS NO.: 200600172631
Improved plant protective efficacy of a baculovirus using an early promoter to drive insect-specific neurotoxin expression 2005

43679 (Item 19 from file: 155) 15422272 PMID: 15133045
Molecular basis of the high insecticidal potency of scorpion alpha-toxins. Jul 23 2004

43680 (Item 20 from file: 5) 0008366569 BIOSIS NO.: 199294068400
MOLECULAR ANALYSIS OF cDNA AND THE TRANSCRIPT ENCODING THE DEPRESSANT INSECT SELECTIVE NEUROTOXIN OF THE SCORPION *LEURUS-QUINQUESTRIATUS -HEBRAEUS* 1992

43681 (Item 21 from file: 5) 0012310167 BIOSIS NO.: 200000028480
Mortality and feeding of mid-stadium larvae of *Helicoverpa zea* and *Helicoverpa armigera* fed a wild strain or a recombinant strain of *Baculovirus* hebraeus expressing an insect-specific toxin of the scorpion *Leurus quinquestriatus hebraeus* 1999

43682 (Item 22 from file: 155) 09708895 PMID: 1801321
Nucleotide sequence and structure analysis of a cDNA encoding an alpha insect toxin from the scorpion *Leurus quinquestriatus hebraeus*. 1991

43683 (Item 23 from file: 5) 0008192856 BIOSIS NO.: 199293035747
NUCLEOTIDE SEQUENCE AND STRUCTURE ANALYSIS OF A cDNA ENCODING AN ALPHA INSECT TOXIN FROM THE SCORPION *LEURUS-QUINQUESTRIATUS -HEBRAEUS* 1991

43684 (Item 24 from file: 155) 14817292 PMID: 12787033
An 'Old World' scorpion beta-toxin that recognizes both insect and mammalian sodium channels. Jun 2003